---Logging off of STN---

=>

Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
FULL ESTIMATED COST	ENTRY 10.68	SESSION 179.19
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY -1.50	SESSION -1.50

STN INTERNATIONAL LOGOFF AT 09:26:03 ON 15 JAN 2006

10/520,426

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID: SSPTAYLC1626

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

```
Web Page URLs for STN Seminar Schedule - N. America
NEWS
NEWS
     2
                 "Ask CAS" for self-help around the clock
NEWS
        DEC 05
                CASREACT(R) - Over 10 million reactions available
     3
NEWS
        DEC 14
                2006 MeSH terms loaded in MEDLINE/LMEDLINE
NEWS
        DEC 14
                2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER
     6 DEC 14
NEWS
                CA/CAplus to be enhanced with updated IPC codes
        DEC 21
NEWS
                IPC search and display fields enhanced in CA/CAplus with the
                 IPC reform
                New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
NEWS
     8
        DEC 23
                 USPAT2
     9
        JAN 13
                 IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS
NEWS 10
         JAN 13
                 New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
                 INPADOC
```

NEWS EXPRESS JANUARY 03 CURRENT VERSION FOR WINDOWS IS V8.01, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005. V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT http://download.cas.org/express/v8.0-Discover/

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NEWS INTER General Internet Information
NEWS LOGIN Welcome Banner and News Items
NEWS PHONE Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

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FILE 'HOME' ENTERED AT 09:23:13 ON 15 JAN 2006

=> file registry
COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 09:23:28 ON 15 JAN 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 13 JAN 2006 HIGHEST RN 871978-06-2 DICTIONARY FILE UPDATES: 13 JAN 2006 HIGHEST RN 871978-06-2

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

Please note that search-term pricing does apply when conducting SmartSELECT searches.

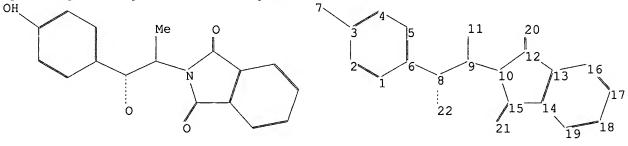
Structure search iteration limits have been increased. See ${\tt HELP\ SLIMITS}$ for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

=>

Uploading C:\Program Files\Stnexp\Queries\10520426\10520426a.str



chain nodes : 7 8 9 11 20 21 22 ring nodes : 1 2 3 4 5 6 10 12 13 14 15 16 17 18 19 chain bonds : 3-7 6-8 8-9 8-22 9-10 9-11 12-20 15-21 ring bonds : 1-2 1-6 2-3 3-4 4-5 5-6 10-12 10-15 12-13 13-14 13-16 14-15 14-19 16-17 17-18 18-19 exact/norm bonds : 3-7 8-22 9-10 10-12 10-15 12-13 12-20 14-15 15-21 exact bonds : 6-8 8-9 9-11 normalized bonds : 1-2 1-6 2-3 3-4 4-5 5-6 13-14 13-16 14-19 16-17 17-18 18-19 isolated ring systems : containing 1 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:Atom 11:CLASS 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom 20:CLASS 21:CLASS 22:CLASS

L1 STRUCTURE UPLOADED

=> d L1 HAS NO ANSWERS L1 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 09:24:02 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 213 TO ITERATE

100.0% PROCESSED 213 ITERATIONS

O ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS:

3385 TO 5135

PROJECTED ANSWERS:

0 TO (

L2 0 SEA

0 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 09:24:17 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 4512 TO ITERATE

100.0% PROCESSED 4512 ITERATIONS

4 ANSWERS

SEARCH TIME: 00.00.01

L3 4 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 167.38 167.59

FULL ESTIMATED COST

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=> d ibib abs hitstr tot
YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y)/N:n

=> file caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.92 168.51

FULL ESTIMATED COST

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FILE COVERS 1907 - 15 Jan 2006 VOL 144 ISS 4 FILE LAST UPDATED: 13 Jan 2006 (20060113/ED)

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=> s 13 L4 2 L3

=> d ibib abs hitstr tot

Same one

W 20030626

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2004:41438 CAPLUS DOCUMENT NUMBER: 140:111274

DOCUMENT NUMBER:

TITLE:

140:111274
Process for preparation of phthalimidophenol
derivatives
Yamaguchi, Toshiaki; Ban, Masaaki; Yanagi, Takashi;
Kikuchi, Ken; Kubota, Minoru; Ozawa, Tetsuji
Kissei Pharmaceutical Co., Ltd., Japan
PCT Int. Appl., 17 pp.
CODEN: PIXXD2
Patent
Japanese
1 INVENTOR (S):

PATENT ASSIGNEE(S): SOURCE:

DOCUMENT TYPE: LANGUAGE:

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

APPLICATION NO. PATENT NO. KIND DATE DATE | MO 2004005251 | AI 20040115 | MO 2003-JP8085 | 20030626 |
| N: AR, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CM, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, KK, MN, MM, MZ, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SZ, SG, SK, SL, SY, TJ, TM, TN, TT, TZ, UA, UG, US, UZ, VC, VN, VY, UZ, AZ, AZ, EW RGH, GM, KE, LS, MM, MZ, SD, SL, SZ, TZ, UG, ZM, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, CG, CM, MI, MR, NE, SN, TD, TG
| CA 2491020 | AA 20040115 | CA 2003-2491020 | 20030626 | EP 1541555 | AI 20050615 | CR 2003-2491020 | 20030626 | RITY APPLN. INFO: | CA 20050928 | US 2005-520426 | 20050107 | RITY APPLN. INFO: | CA 20030928 | AI 20050922 | US 2005-520426 | 20050107 | CA 20030626 | AI 20050928 | US 2005-520426 | 20050107 | AI 20030626 | AI 20050928 | US 2005-520426 | 20050107 | AI 20030626 | AI 20050928 | US 2005-520426 | 20050107 | AI 20030626 | AI 2 PRIORITY APPLN. INFO .:

GI

. . . .

L4 ANSWER 2 OF 2
ACCESSION NUMBER:
DSGULMENT NUMBER:
171TLE:
AUTHOR(S):
CORPORATE SOURCE:
Tech. Hochach. Otto Von Guericke, Magdeburg, Fed.

WO 2003-JP8085

Rep.

Ger.

Archiv der Pharmazie und Berichte der Deutschen Pharmazeutischen Gesellschaft (1968), 301(2), 161-7 CODEN: APBDAJ; ISSN: 0376-0367

DOCUMENT TYPE:

LANGUAGE:

Gernan

OTHER SOURCE(S):

Gernan

OTHER SOURCE(S):

Gernan

Nah in C6H6 1hr. under reflux, followed by saponification with 0.3 g. p-MecGH4SO3H to give 42.5% I (R = EL) (II), m. 111-13'. II (6.93 g.) was brominated to give 84% I (R = CKBRMP) (III), m. 110-12'.

On treatment with K phthalimide at 50' III formed 26.2% I (R = (a-phthalimid) -thyl), m. 130-1'. IV was heated to 210' 4 hrs. with pyridine hydrochloride to give 91.5% p-MecGH4COCHR'Me (R' = phthalimid), m. 180-1', which was nitrated at room temperature to give 73% 3,5,4-(02N)2(HO)CGH2COCHR'Me (VI), m. 165-6'. On treatment with PMOCI3 V gave 99% 3,54-(02N)2CIGGH2COCHR'Me (VI), m. 192-3.5'. VI was condensed with p-MecGH4COCHGHMe (VI), m. 180-1'. Was condensed with p-MecGH4COCHGHMe (VI), m. 180-1', which was nitrated at room temperature to give 73% 3,5,4-(02N)2(HO)CGH2COCHR'Me (VI), m. 165-6'. On treatment with PMOCI3 V gave 99% 3,54-(02N)2CIGGH2COCHR'Me (VI), m. 192-3.5'. VI was condensed with p-MecGH4COCHGH4H in pyridine 48 hrs. to give 90% I [R-(a-phthalimido)ethyl) (VIII), m. 134-6'. On reduction with PMOC followed by diazotization and treatment with iddine and NaI, VII gave 30% 3,54-12(p-MeoC6H4O)CGH2COCHR'Me, m. 188-8.5', which was

30% 3.5.4-I2(p-MeOCSH40)C6H2COCHR'Me, m. 188-8.5*, which was reduced with iso-PrON and (iso-PrO)3Al to give 67% 3.5.4-I2(p-MeOC6H40)C6H2COHR'Me m. 225-6*. Treatment of 3.5.4 (102N)2 (HO)C6H4COCHR'Me or 4-(102N)2C1C6H4COCHR'Me with hydroquinone gave 3.5.4-(102N)2 (p-HOC6H40)C6H2COCHR'Me (VIII) m. 204-5* in 76.3 and 71.38 yield, resp. VIII was converted to 32% 3.5.4-12(p-HOC6H40)C6H2COCHR'Me, m. 213-15*, by reduction, diazotization and treatment with iodine and Na1. 19427-12-4P 19427-13-5P RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of) 19427-12-4 CAPLUS Phthalimide, N-(p-hydroxy-α-methylphenacyl)-, DL- (BCI) (CA INDEX NAME)

IT

19427-13-5 CAPLUS

Phthalimide, N-(4-hydroxy-α-methyl-3,5-dinitrophenacyl)-, DL- (8CI) (CA INDEX NAME)

ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)
This invention pertains to a method for producing novel phenol derivs.
with formula of I, II, and III. For example, I was prepared starting from

(S)-2-phthalimidopropionic acid and anisole in PhCl in the presence of SOC12 and AlC13 in 53.6% yield with 99.9% e.e. I was hydrogenated with

in MeOH in the presence of 10% Pd/C to give II (57%) with 100% e.e. II was nydrogenated with was treated with MeNH2 in MeoH to afford III (54%) with 100% e.e. The compds. I-III are useful as intermediates for various medicines. 646037-20-9P 646037-21-0P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation of phthalimidophenol derivs.) 646037-20-9 CAPLUS IM-Isoindole-1,3(2H)-dione, 2-[(15)-2-(4-hydroxyphenyl)-1-methyl-2-oxoethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

646037-21-0 CAPLUS 1H-Isoindole-1,3(2H)-dione, 2-[(15,2R)-2-hydroxy-2-(4-hydroxyphenyl)-1-methylethyl]- (9C1) (CA INDEX NAME)

Absolute stereochemistry.

REFERENCE COUNT:

FORMAT

21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR

RECORD. ALL CITATIONS AVAILABLE IN THE RE

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)